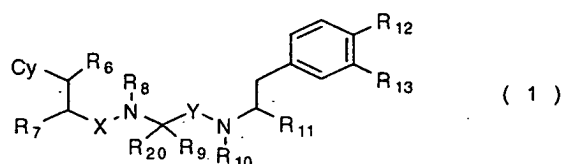


Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

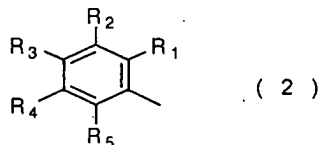
Listing of Claims:

1. (Currently Amended) A compound of Formula (1):



wherein:

Cy is a group of Formula (2):



~~C₃-cycloalkyl or phenyl,~~

~~R₁, R₂, R₃, R₄ and R₅ are hydrogen, halogen, or hydroxy, amino, trifluoromethyl or nitrile and at least one of R₁, R₂, R₃, R₄ and R₅ is halogen, trifluoromethyl or nitrile;~~

~~R₆ is hydrogen, optionally substituted straight-chained or branched C₁₋₃ alkyl, amino or hydroxy;~~

~~R₇ is hydrogen, optionally substituted straight-chained or branched C₁₋₃alkyl, substituted with one or more hydroxyl groups, or amino optionally substituted with one or~~

~~more of the same or different kinds of straight-chained or
branched C₁₋₃ alkyl groups which may be the same or different,
or hydroxy;~~

R₈ is hydrogen, methyl or ethyl;

R₉ is ~~optionally substituted~~ straight-chained or
branched C₁₋₆ alkyl optionally substituted with one or more
groups which may be the same or different and are selected
from the group consisting of phenyl, para-hydroxyphenyl, para-
fluorophenyl, para-chlorophenyl, C₃₋₇ cycloalkyl, halogen and
thienyl, ~~optionally substituted straight chained or branched
C₂₋₆alkenyl, optionally substituted straight chained or
branched C₂₋₆alkynyl, C₃₋₇cycloalkyl; or optionally substituted
phenyl;~~

R₂₀ is ~~hydrogen or straight chained or branched
C₁₋₃alkyl or R₉ and R₂₀ may together form C₃₋₇cycloalkyl;~~

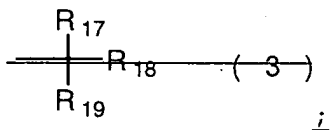
R₁₀ is hydrogen or methyl or ethyl ~~straight chained
or branched C₁₋₃alkyl;~~

R₁₁ is ~~hydrogen,~~ straight-chained or branched C₁₋₃
alkyl optionally substituted with one or more groups which may
be the same or different and are selected from the group
consisting of ~~amino optionally substituted with one or more of
the same or different straight chained or branched C₁₋₃ alkyl,
3 to 7 membered cyclic amino optionally substituted with
hydroxyl, amino, carboxyl, carbamoyl or methyl; hydroxyl,~~

~~methoxy, halogen, carbamoyl, methanesulfonyl, ureide,~~
~~guanidyl, N'-cyano-N"-methylguanidyl, sulfamoylamino,~~
~~carbamoylmethylamino and methanesulfonylamino, and -CO-~~
~~N(R₁₄)R₁₅-carboxyl;~~

~~R₁₂ is hydroxy or OR₁₆;~~

~~R₁₃ is hydrogen, straight-chained or branched C₁₋₆~~
~~alkyl, straight-chained or branched C₂₋₆alkenyl, straight-~~
~~chained or branched C₂₋₆alkynyl or a group of Formula (3):~~



~~R₁₄ and R₁₅, which may be the same or different, are~~
~~each hydrogen, straight-chained or branched C₁₋₃ alkyl~~
~~optionally substituted with straight-chained or branched C₁₋₃~~
~~alkoxy optionally substituted with hydroxyl, amino, carboxyl~~
~~or carbamoyl, hydroxyl, amino, methylamino, dimethylamino,~~
~~carbamoyl or methanesulfonyl; optionally substituted straight-~~
~~chained or branched C₁₋₄ alkyl, C₃₋₇cycloalkyl, straight-chained~~
~~or branched C₁₋₄ alkoxy, straight-chained or branched C₁₋₄~~
~~alkylsulfonyl, or pyridyl; a heterocyclic ring;~~

~~R₁₆ is straight-chained C₁₋₄ alkyl;~~

~~R₁₇ is hydrogen or methyl;~~

~~R₁₈ and R₁₉ together form cycloalkyl or C₃₋₇~~
~~cycloalkenyl;~~

X is carbonyl or methylene;

Y is carbonyl ~~or~~ methylene;

or a pharmaceutically acceptable salt thereof.

2. (Previously presented) The compound according to claim 1,
wherein Cy in Formula (1) is a group of Formula (2);
or a pharmaceutically acceptable salt thereof.

3. (Previously presented) The compound according to claim 1,
wherein Cy in Formula (1) is a group of Formula (2) in which
at least one of R₁, R₂, R₃, R₄ and R₅ is halogen and the others
are hydrogen or hydroxy;
or a pharmaceutically acceptable salt thereof.

4. (Previously presented) The compound according to claim 1,
wherein Cy in Formula (1) is a group of Formula (2) in which
R₃ is halogen or R₂ and R₃ are the same kind of halogen;
or a pharmaceutically acceptable salt thereof.

5. (Previously presented) The compound according to claim 1,

wherein Cy in Formula (1) is a group of Formula (2) in which R₃ is halogen and R₁, R₂, R₄ and R₅ are hydrogen, or R₂ and R₃ are the same kind of halogen and R₁, R₄ and R₅ are hydrogen; or a pharmaceutically acceptable salt thereof.

Claims 6-13. (Canceled)

14. (Previously presented) The compound according to claim 1, wherein R₇ in Formula (1) is hydrogen or amino optionally substituted with one or more of the same of different kinds of straight-chained or branched C₁₋₃ alkyl; or a pharmaceutically acceptable salt thereof.

15. (Previously presented) The compound according to claim 1, wherein R₈ in Formula (1) is hydrogen or methyl; or a pharmaceutically acceptable salt thereof.

16. (Previously presented) The compound according to claim 1, wherein R₉ in Formula (1) is methyl, isopropyl, isobutyl, sec-butyl, tert-butyl, 3-pentyl, neopentyl, cyclohexyl, phenyl, benzyl, para-hydroxybenzyl, cyclohexylmethyl or para-fluorobenzyl; or a pharmaceutically acceptable salt thereof.

Claims 17-18. (Cancelled)

19. (Currently Amended) The compound according to claim 1, wherein R₁₁ in Formula (1) is methyl, hydroxymethyl,

carbamoylmethyl, methanesulfonylmethyl, ureidemethyl,
sulfamoylaminomethyl, methanesulfonylaminomethyl, ~~carbamoyl,~~
ethylcarbamoyl, n-propylcarbamoyl, isopropylcarbamoyl,
~~cyclopropylcarbamoyl,~~ tertbutylcarbamoyl, methoxycarbamoyl,
methylcarbamoyl, methanesulfonylmethylcarbamoyl,
methoxymethylcarbamoyl,;

or a pharmaceutically acceptable salt thereof.

Claim 20 Cancelled

21. (Currently Amended) The compound according to
claim 1, wherein R_{13} in Formula (1) is isopropyl, tert-butyl
(tBu), or 1,1-dimethylpropyl ~~or 1,1-dimethyl-2-propenyl;~~
or a ~~hydrate or~~ pharmaceutically acceptable salt thereof.

22. (Currently Amended) The compound according to
claim 1, wherein in Formula (1) Cy is a group of Formula (2)
in which at least one of R_1 , R_2 , R_3 , R_4 and R_5 is halogen and
the others are hydrogen or hydroxy;

~~R_6 is hydrogen or methyl;~~

~~R_7 is hydrogen or optionally substituted amino optionally
substituted with one or more of the same or different
straight chained or branched C_{1-3} alkyl;~~

R_8 is hydrogen or methyl;

R₉ is methyl, isopropyl, isobutyl, sec-butyl, tert-butyl, 3-pentyl, neopentyl, cyclohexyl, phenyl, ~~benzyl, para-hydroxybenzyl, para-fluorobenzyl or cyclohexylmethyl;~~
~~R₁₀ is hydrogen;~~
~~R₁₁ is hydrogen or methyl;~~
R₁₁ is methyl, hydroxymethyl, carbamoylmethyl, methanesulfonylmethyl, ureidemethyl, sulfamoylaminomethyl, methanesulfonylaminomethyl, ~~carbamoyl,~~ methylcarbamoyl, ethylcarbamoyl, n-propylcarbamoyl, isopropylcarbamoyl, ~~tert-cyclopropylcarbamoyl butylcarbamoyl,~~ , methanesulfonylmethylcarbamoyl, methoxymethylcarbamoyl, or methoxycarbamoyl;
~~R₁₂ is hydroxy;~~
R₁₃ is isopropyl, tert-butyl (tBu), 1,1-dimethylpropyl-or 1,1-dimethyl-2-propenyl;
or a pharmaceutically acceptable salt thereof.

23. (Previously presented) The compound according to claim 1 which is selected from the group of compounds consisting of Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, Phe(4-Cl)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, Phe(3,4-F₂)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, Phe(3-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHOMe, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-tertbutyl-4-hydroxyphenyl)-1-(2-

pyridylcarbamoyl)ethylamide, N-(2-(2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methyl-butyrylamino)-3-(3-tBu-4-hydroxyphenyl)propyl)urea, N-(2-(2-(2-amino-3-(4-fluorophenyl)propanoyl-N-methylamino)-3-methyl)butyrylamino)-3-(3-tertbutyl-4-hydroxyphenyl)propyl)sulfamide, N-[2-(3-tertbutyl-4-hydroxyphenyl)-1-(methanesulfonylaminomethyl)ethyl]-2-[N-(4-fluorophenylalanyloyl)methylamino]-3-methylbutanamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-carbamidemethylethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-methanesulfonylmethylethylamide, 2-(2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methyl-butyrylamino)-3-(3-tBu-4-hydroxyphenyl)propanol, 2-(1-(2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methyl-butyrylamino)-2-(3-tertbutyl-4-hydroxyphenyl)ethyl)-6-methyl-4-pyrimidinone, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-(1,3,4-oxadiazol-2-yl)ethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-(1,2,4-oxadiazol-5-yl)ethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-

methylbutyric acid 2-(3-tertbutyl-4-hydroxyphenyl)-1-(thiazol-2-yl)ethylamide, 2-((2-amino-3-(4-fluorophenyl)propionyl)-N-methylamino)-3-methylbutyric acid 2-(3-t-butyl-4-hydroxyphenyl)-1-(1,3,4-triazol-2-yl)ethylamide, Tyr(2-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, Tyr(3-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH₂, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH₂, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH₂, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHMe, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, N-Et-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHMe, N-Et-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHMe, Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH₂, N-Me-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH₂, N-Et-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH₂, Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NHMe, N-Me-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NHMe, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHtBu, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHCH₂SO₂CH₃, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH₂Et, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH₂Et, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NH₂Et, Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHCH₂OH, N-Me-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHCH₂OH, N-Et-Phe(4-F)-N-Me-Val-Tyr(3-tBu)-NHCH₂OH, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂Et, N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂Et, N-Et-Phe(4-F)-N-Me-Val-N-Me-

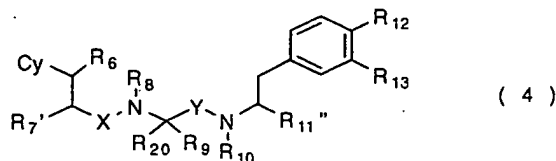
Tyr(3-tBu)-NH₂Et, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHCH₂OH,
N-Me-Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NHCH₂OH, N-Et-Phe(4-F)-
N-Me-Val-N-Me-Tyr(3-tBu)-NHCH₂OH, Phe(4-F)-N-Me-Val-N-Et-
Tyr(3-tBu)-NH₂Et, N-Me-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH₂Et,
N-Et-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-NH₂Et, Phe(4-F)-N-Me-
Val-N-Et-Tyr(3-tBu)-NHCH₂OH, N-Me-Phe(4-F)-N-Me-Val-N-Et-
Tyr(3-tBu)-NHCH₂OH, N-Et-Phe(4-F)-N-Me-Val-N-Et-Tyr(3-tBu)-
NHCH₂OH, Phe(4-F)-N-Me-Val-N-Me-Tyr(3-tBu)-NH₂Pr, and Phe(4-
F)-N-Me-Val-Tyr(3-tBu)-NH₂Pr Phe(4-F)-N-Me-Val-Tyr(3-tBu)-
NH₂Pr;
or a pharmaceutically acceptable salt thereof.

24. (Previously Presented) A pharmaceutical composition containing an effective amount of the compound according to claim 1 as an active ingredient and an inert pharmaceutically acceptable carrier.

25. (Previously Presented) A motilin receptor antagonist composition containing an effective amount of the compound according to claim 1 and an inert pharmaceutically acceptable carrier.

Claims 26-27. (Cancelled)

28. (Currently Amended) A compound of Formula (4):



wherein

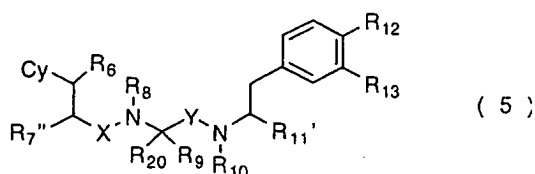
Cy, R₆, R₈, R₉, R₂₀, R₁₀, R₁₂, R₁₃, X and Y are as defined in claim 1;

R₇' is ~~hydrogen, straight-chained or branched C₁₋₃ alkyl substituted with one or more optionally having at least one-protected hydroxyl groups substituent, or protected amino optionally substituted with having at least one or more substituent of the same or different straight-chained or branched C₁₋₃ alkyl groups which may be the same or different or protected hydroxyl; and~~

R₁₁'' is ~~hydrogen, straight-chained or branched C₁₋₃ alkyl optionally substituted with one or more groups which may be the same or different and are selected from the group consisting of amino optionally substituted with one or more of the same or different straight chained or branched C₁₋₃ alkyl, 3 to 7 membered cyclic amino optionally substituted with hydroxyl, amino, carboxyl, carbamoyl or methyl, hydroxyl, methoxy, halogen; carbamoyl, methanesulfonyl, ureide, guanidyl, N'-cyano-N''-methylguanidyl, sulfamoylamino, carbamoylmethylamino, and -CO-~~

$N(R_{14})R_{15}$, wherein R_{14} and R_{15} are as defined in claim 1,
~~carboxyl, straight chained or branched C_{1-3} alkyl having a~~
~~protected amine;~~
or a pharmaceutically acceptable salt thereof.

29. (Currently Amended) A compound of Formula (5):



wherein:

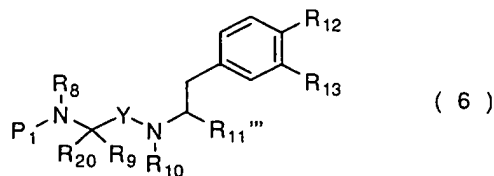
Cy , R_6 , R_8 , R_9 , R_{10} , R_{12} , R_{13} , X and Y are as defined in claim 1;

R_7 is ~~hydrogen,~~ straight-chained or branched C_{1-3} alkyl ~~optionally having at least optionally substituted with~~
~~one or more optionally protected hydroxyl groups substituent~~
~~or amino optionally having at least substituted with one or~~
~~more substituents which are the same or different straight-~~
~~chained or branched C_{1-3} alkyl groups which may be the same or~~
~~different, or optionally protected hydroxy;~~ and

R_{11}' is ~~hydrogen,~~ straight-chained or branched C_{1-3} alkyl ~~optionally substituted with one or more groups having~~
~~at least one protected substituent s~~ which may be the same or
different and are selected from the group consisting of
~~protected amino optionally substituted with one or more~~

~~straight-chained or branched C₁₋₃ alkyl, protected 3 to 7~~
~~membered cyclic amino optionally substituted with protected~~
~~hydroxyl, protected amino, protected carboxyl or protected~~
~~carbamoyl; protected hydroxyl₁, protected carbamoyl₁,~~
~~protected ureide₁, protected guanidyl₁, protected N'-cyano-~~
~~N''-methylguanidyl₁, protected sulfamoylamino₁, protected~~
~~carbamoylmethylamino and protected methanesulfonylamino; and~~
~~-CO-N(R₁₄)R₁₅ wherein R₁₄ and R₁₅ are as those defined in claim 1~~
~~which are appropriately protected, carboxyl~~
or a ~~hydrate or~~ pharmaceutically acceptable salt thereof.

30. (Currently Amended) A compound of Formula (6):



wherein:

R₈ is hydrogen, methyl or ethyl~~optionally~~
~~substituted straight chained or branched C₁₋₃ alkyl, optionally~~
~~substituted amino, or hydroxy;~~

R₉, is ~~optionally substituted~~ straight-chained or
branched C₁₋₆ alkyl optionally substituted with one or more
groups which may be the same or different and are selected
from the group consisting of phenyl, para-hydroxyphenyl, para-
fluorophenyl, para-chlorophenyl, C₃₋₇ cycloalkyl, halogen and

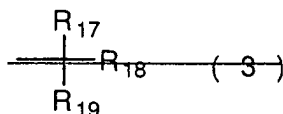
~~thienyl, optionally substituted straight chained or branched~~
~~C₂₋₆ alkenyl, optionally substituted straight chained or~~
~~branched C₂₋₆ alkynyl, C₃₋₇ cycloalkyl or optionally substituted~~
~~phenyl;~~

R₂₀ is hydrogen or methyl ~~or straight chained or~~
~~branched C₁₋₃ alkyl, or R₉ and R₂₀ may together form C₃₋₇~~
~~cycloalkyl;~~

R₁₀ is hydrogen or methyl or ethyl ~~straight chain or~~
~~branched C₁₋₃ alkyl;~~

R₁₂ is hydroxy ~~or OR₁₆;~~

R₁₃ is ~~hydrogen,~~ straight-chained or branched C₁₋₆
~~alkyl, straight chained or branched C₂₋₆ alkenyl, straight~~
~~chained or branched C₂₋₆ alkynyl or a group of Formula (3)~~



~~Wherein R₁₇ is hydrogen or methyl,~~

~~R₁₈ and R₁₉ together form cycloalkenyl or C₃₋₇~~
~~cycloalkenyl; and~~

Y is carbonyl ~~or methylene;~~

P₁ is hydrogen or a protecting group of amine; and

R_{11'''} is ~~hydrogen,~~ straight-chained or branched C₁₋₃alkyl,
carboxyl, straight-chained or branched C₁₋₃alkyl optionally
substituted with one or more groups which may be the same or
different and are selected from the group consisting of amino

~~optionally substituted with one or more of the same or~~
~~different straight chained or branched C₁₋₃ alkyl, 3 to 7~~
~~membered cyclic amino optionally substituted with hydroxyl,~~
~~amino, carboxyl, carbamoyl or methyl, hydroxyl, methoxy,~~
~~halogen, carbamoyl, methanesulfonyl, ureide, guanidyl,~~
~~N'-cyano-N''-methylguanidyl, sulfamoylamino,~~
~~carbamoylmethylamino and methanesulfonylamino; carboxyl,~~
straight-chained or branched C₁₋₃ alkyl having protected amino
or an ~~optionally substituted heterocyclic ring, or and~~ -CO-
N(R₁₄)R₁₅ wherein R₁₄ and R₁₅, which may be the same or
different, are hydrogen, ~~optionally substituted straight-~~
chained or branched C₁₋₄ alkyl optionally substituted with
hydroxy, C₃₋₇ cycloalkyl, straight-chained or branched C₁₋₄
alkoxy, straight-chained or branched C₁₋₄alkylsulfonyl, or
pyridyl ~~a heterocyclic ring, carboxyl, straight chained or~~
~~branched C₁₋₃alkyl having protected amino or an optionally~~
~~substituted heterocyclic ring;~~
or a pharmaceutically acceptable salt thereof.

Claims 31-34. (Canceled)

35. (Previously Presented) The compound according
to claim 1, wherein the substitution of the optionally
substituted straight-chained or branched C₁₋₃ alkyl as R₇ in
formula (1) is halogen, hydroxyl or amino.